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END OF FISCAL YEAR REPORT

OCTOBER 1, 1983 - SEPTEMBER 30, 1984

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CHARACTERIZATION OF NOVEL BINDERS BASED ON ENERGETIC THERMOPLASTIC ELASTOMERS

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17. FIELD	GROUP	SUB. GR			(Continue on reverse if necessary and identify by block number) A. Energetic Thermoplastic Elastomers, PolyBEMO,					
*		Block copolymers, polyethers					. rorysemo,			
19. ABSTR	ACT (Continue	on reverse if nece	ssory and	l identify by block numbe	r)					
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Dr. Richard S. Hiller

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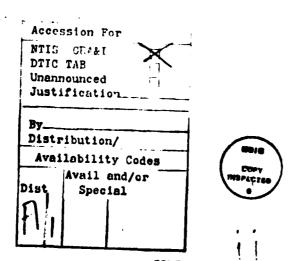
SECURITY CLASSIFICATION OF THIS PAGE

- 4. Processing can be carried out in the range of 80-110°C.
- 5. PolyBEMO and polyBAMO are highly crystalline materials. PolyBEMO exhibits more than one crystalline form.

The major focus of the research has been to characterize the thermal and physical behavior of these materials. While the work is still incomplete, major accomplishments of the past year are as follows:

- 1. Thermal degradation studies on poly[3,3-bis(ethoxy methyl) oxetane], polyBEMO. This research utilized both differential scanning calorimetry, DSC, and mass spectrometry, showing that C-O bonds in both the backbone and side chains were the main source of degradation during heating. This reaction was demonstrated to be endothermic in nature, important in preventing detonation of the propellent by hot spall.
- 2. In a basic characterization study of di- and triblock copolymers based on polyBEMO, the melting temperature, T_m, of polyBEMO was found to be 85°C, while its glass transition temperature was found to be about -30°C. In a triblock copolymer with a random copolymer center block of poly(BNMO-co-THF), the modulus was found to be in the leathery range through the proposed application temperatures. Crystallization was found to be complete after 2-5 minutes. In terms of processing temperature range and general physical properties, these materials were found to be outstanding candidates for LOVA binder materials.
- 3. In the most recent study, the nature of crystallinity in polyBENO and poly[3,3-bis(azidomethyl) oxetane], polyBANO was examined. PolyBENO was found to be the more crystalline, while the heat of crystallization is larger for polyBANO. Two crystalline forms were identified by x-ray studies for polyBENO, while only one crystalline form was found for polyBENO. The thermodynamics of fusion were investigated via DSC and melting point depression studies, providing important energy requirements for future manufacturing needs.

Recently, much interest has been expressed on the development of thermoplastic elastomers with crystalline hard blocks, rather than glassy blocks. The main advantages of such materials include the possibility of having the two blocks being miscible in the melt, with the hard block crystallizing out on cooling. Such a system would have lover melt viscosities than corresponding products exhibiting phase separation in the melt. The present systems are giving good evidence that this may be true in important materials of LOVA interest. We think that the block copolymers based on polyBEMO, polyBAMO, and related material offer much promise in the development of novel elastomers, adhesives, and binders. As already mentioned, these materials also have melting temperatures and other properties that are known to be satisfactory.



LIST OF PRESENTATIONS:

- "Polyether Block Copolymers for the LOVA Program", ONR Workshop, U. Mass., MA, May 1983.
- 2. "Thermodynamics of Phase Separation in Crystalline Block Copolymers Based on Polyethers", Capri, Italy, International Symposium of Phase Relationships and Properties of Multicomponent Polymer Systems", May 1983.
- 3. "Characterization of Block Copolymers Based on PolyBEMO", ONR Workshop, Chesapeake Bay Meeting, July, 1983.
- 4. "Characterization of Novel Polyethers for the LOVA Program", ONR Workshop, Chestertown, MD, August, 1982.
- 5. "Viscoelastic Behavior of Block Copolymers Based on Polyethers", AIChE Meeting, Los Angeles, CA, November, 1982.

LIST OF PAPERS AND PUBLICATIONS:

- 1. R. B. Jones, C. J. Murphy, L. H. Sperling, M. Farber, S. P. Harris, and G.E. Manser, "Thermal Decomposition Behavior of Poly[3,3-bis(ethoxymethyl) oxetane] and Related Polyethers", accepted, Journal of Applied Polymer Science.
- 2. K. E. Hardenstine, C. J. Murphy, R. B. Jones, and G. E. Manser, "Characterization of Block Copolymers Based on Poly[3,3-bis(ethoxymethyl) oxetane] and other Novel Polyethers", accepted, Journal of Applied Polymer Science.
- 3. K. E. Hardenstine, G. V. Henderson, Jr., L. H. Sperling, C. J. Murphy, and G. E. Manser, "Crystallization Behavior of Poly[3,3-bis(ethoxymethyl) oxetane] and Poly[3,3-bis(azidomethyl oxetane], submitted, J. Polymer Science, Polymer Physics Edition.

L. H. Sperling

Sources of Support Fiscal Year 1984 (in addition to ONR 533613)

	Agency		Annual Support \$
1.	nsf	DMR-8405053 "Structure and Morphology of IPN's Using Small-Angle Neutron Scattering and Phase Diagram Methods"	\$66,000
2.	NSF	CPE-8206720 "Neutron Scattering from Latex Particles"	\$66,000
3.	U.S. Navy	N00014-84-K-0508 "The Fundamental Nature of the Damping Phenomenon as Characterized with Tailored IPN Compositions"	\$80,000

Personnel

- Dr. L. H. Sperling, Principal Investigator
- Dr. C. J. Murphy, Visiting Scientist
- Ms. K. E. Hardenstine, M.S. Candidate, graduated October, 1984.
- Mr. R. B. Jones, M.S. Candidate, graduated June, 1984.
- Mr. G. V. Henderson, M.S. Candidate

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